

Sub A

Tryptase  $\alpha$  MSLSLLLALP VLASRAYAAP APVQALQQAG IVGGQEAPRS KWPWQVSLRV RDRYWMHFCG (30)

Tryptase  $\beta$ -I --N----- --G---RV- ----- HGP-----

Tryptase  $\beta$ -II --N----- --G---RV- ----- HGP-----

A

Tryptase  $\alpha$  GSLIHPQWVL TAAHCLGPDV KDLATLRVQL REQHLVYQDQ LLPVSRIIVH PQFYFIQTGA (90)

Tryptase  $\beta$ -I -----V-----A----- --TA-I--

Tryptase  $\beta$ -II -----V-----A----- --TA-I--

B

C

Tryptase  $\alpha$  DIALLELEEP VNISSRVHTV MLPPASETFP PGMPCWVTGW GDVDNDEPLP PPFPKQVKV (150)

Tryptase  $\beta$ -I -----V-H--- T-----R-----

Tryptase  $\beta$ -II -----KV-H--- T-----R-----

D

Tryptase  $\alpha$  PIMENHICDA KYHLGAYTGD DVRIIRDML CAGNSQRDSC KGDSGGPLVC KVNGLWLQAG (210)

Tryptase  $\beta$ -I -----V-----TR--- Q-----

Tryptase  $\beta$ -II -----V-----TR--- Q-----

3

1

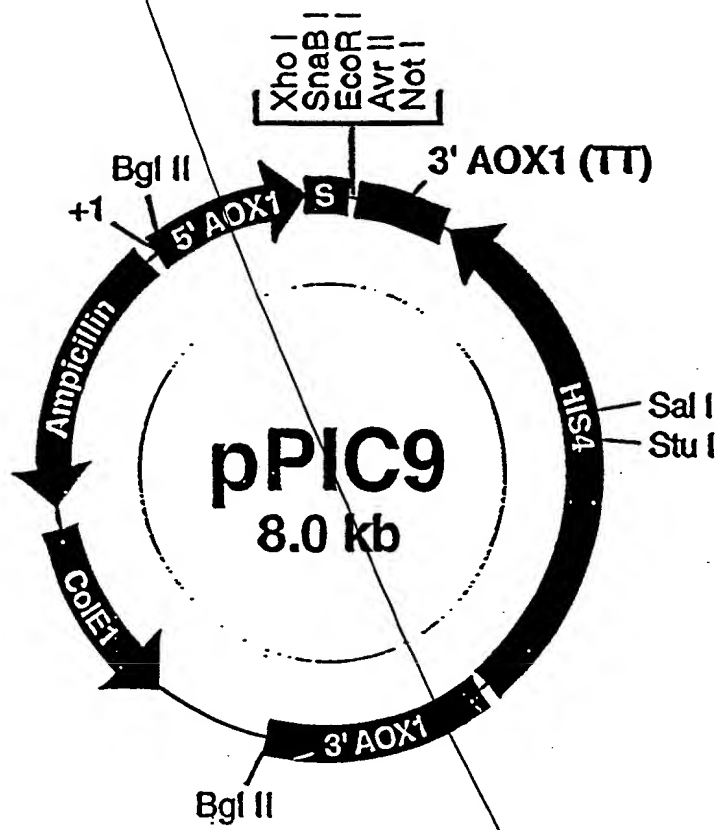
Tryptase  $\alpha$  VVSWDEGCAQ PNRPGIYTRV TYILDWIIHY VPKKP

Tryptase  $\beta$ -I -----G-----

Tryptase  $\beta$ -II -----G-----

2

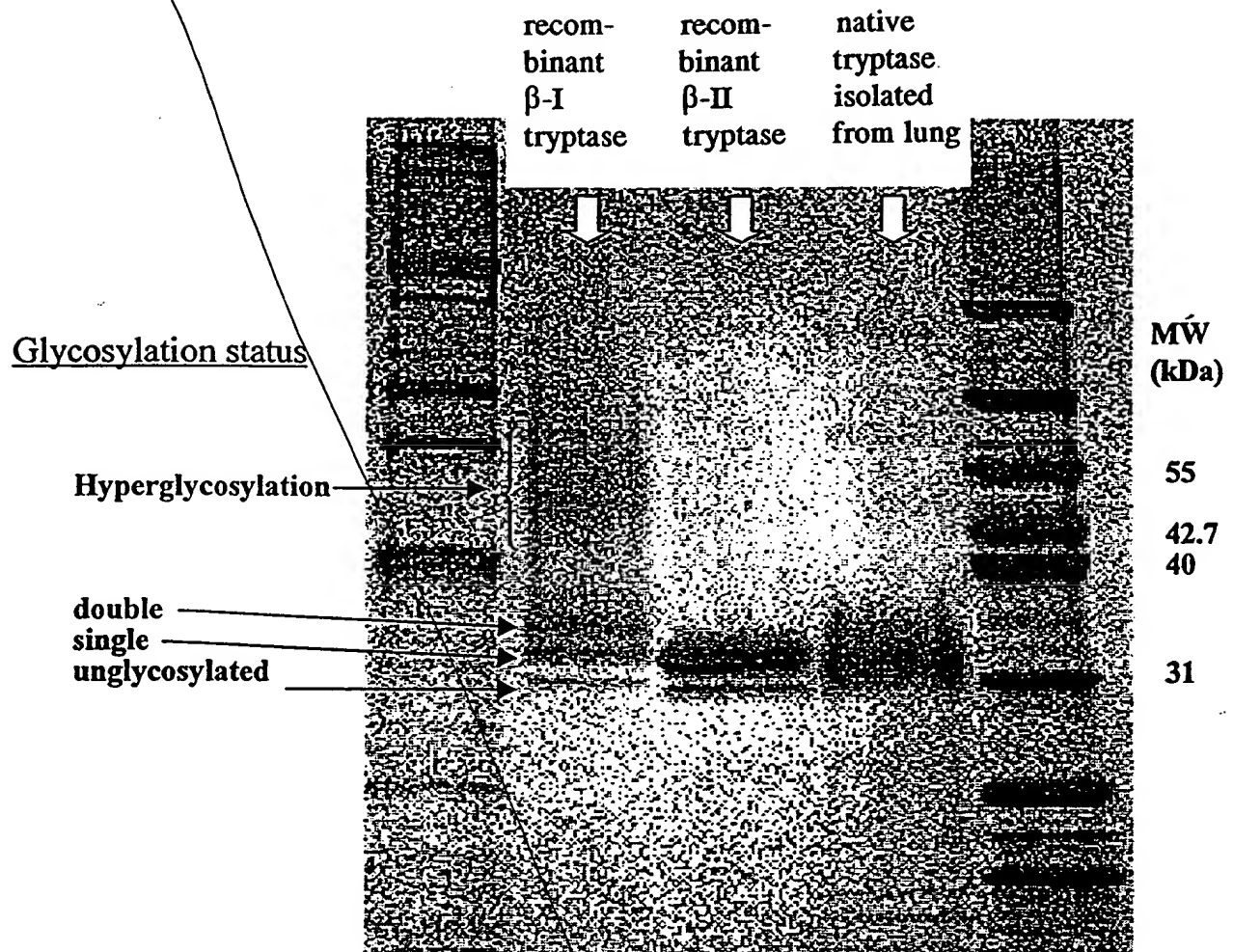
FIG. 1



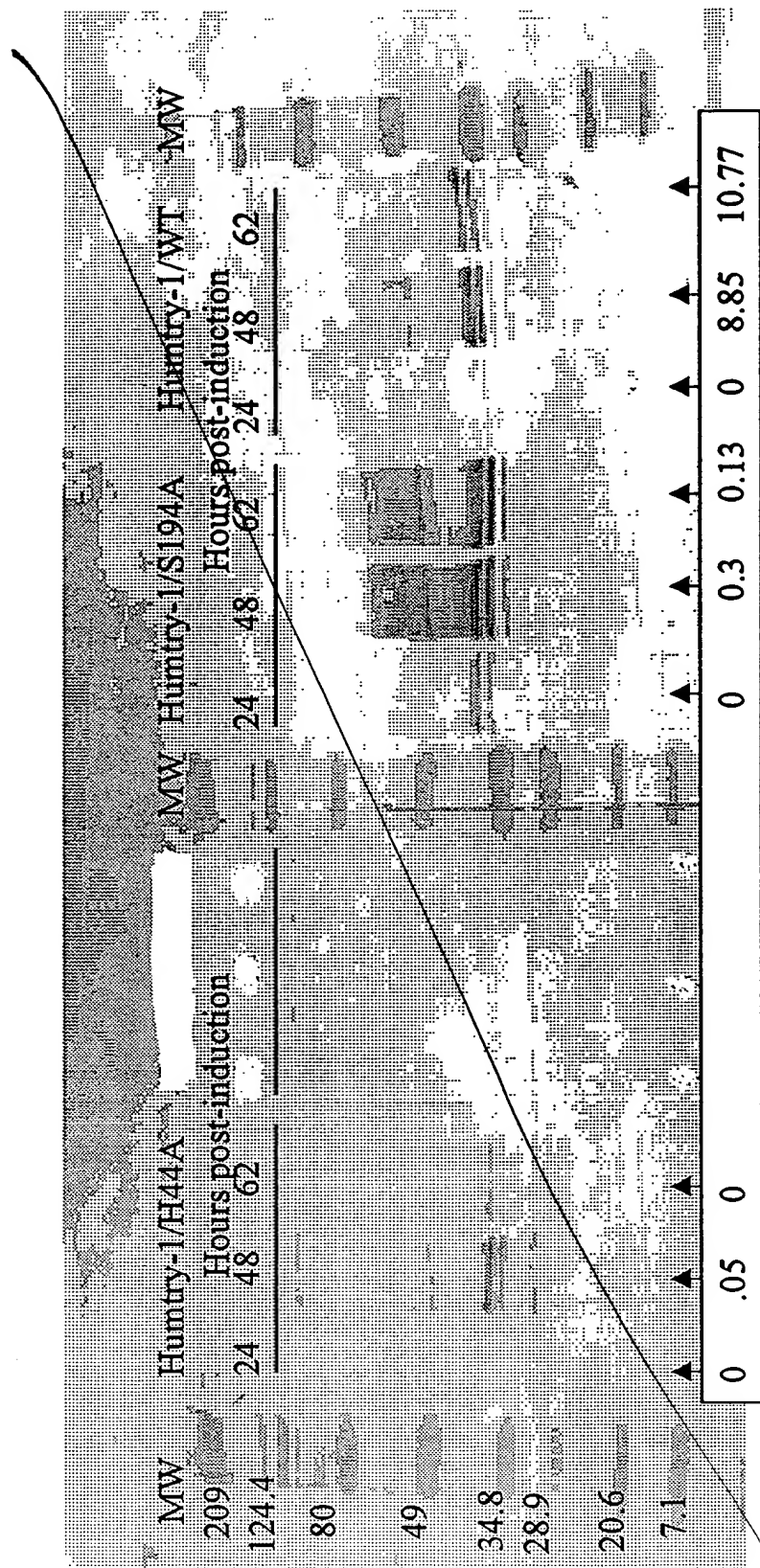
**FIG. 2**

SDS-PAGE gel showing the digestion of rhβI and rhβII with Native β. The gel is divided into two main sections: 'Undigested' and 'Digested'. Each section has lanes for Native β, rhβI, and rhβII. Molecular weight markers are shown on the right at 34 kD and 29 kD. Arrows indicate the positions of the bands.

**FIG. 3**

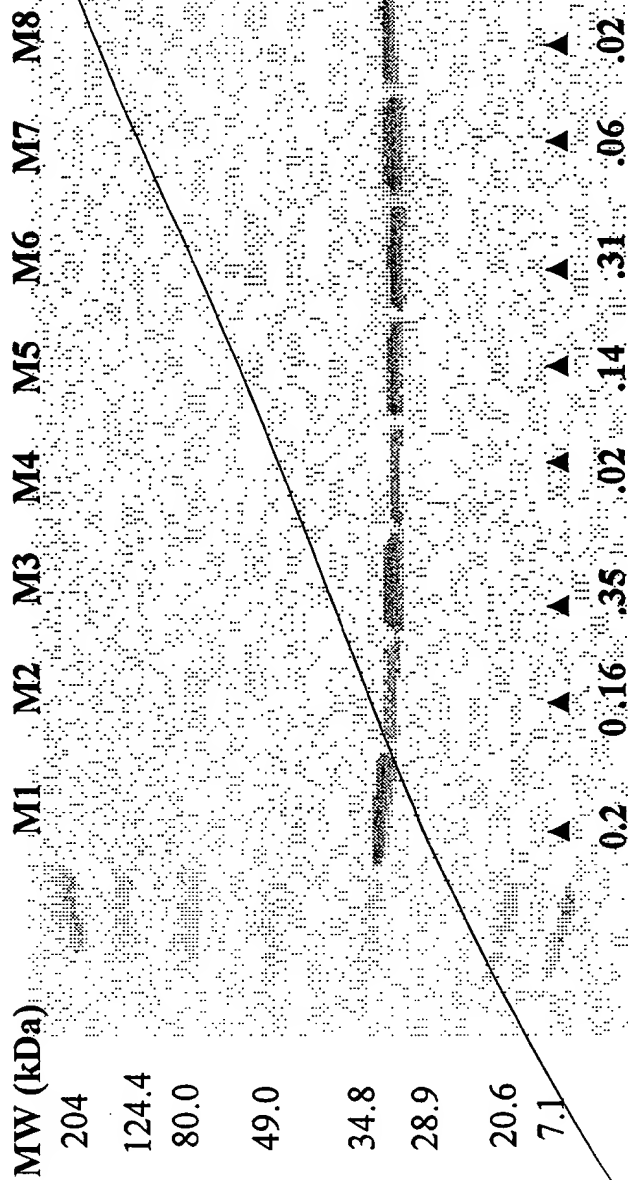


**FIG. 4**



**FIG. 5**

[-----Humtry-1/N102K/S194A Mutant Clones-----]

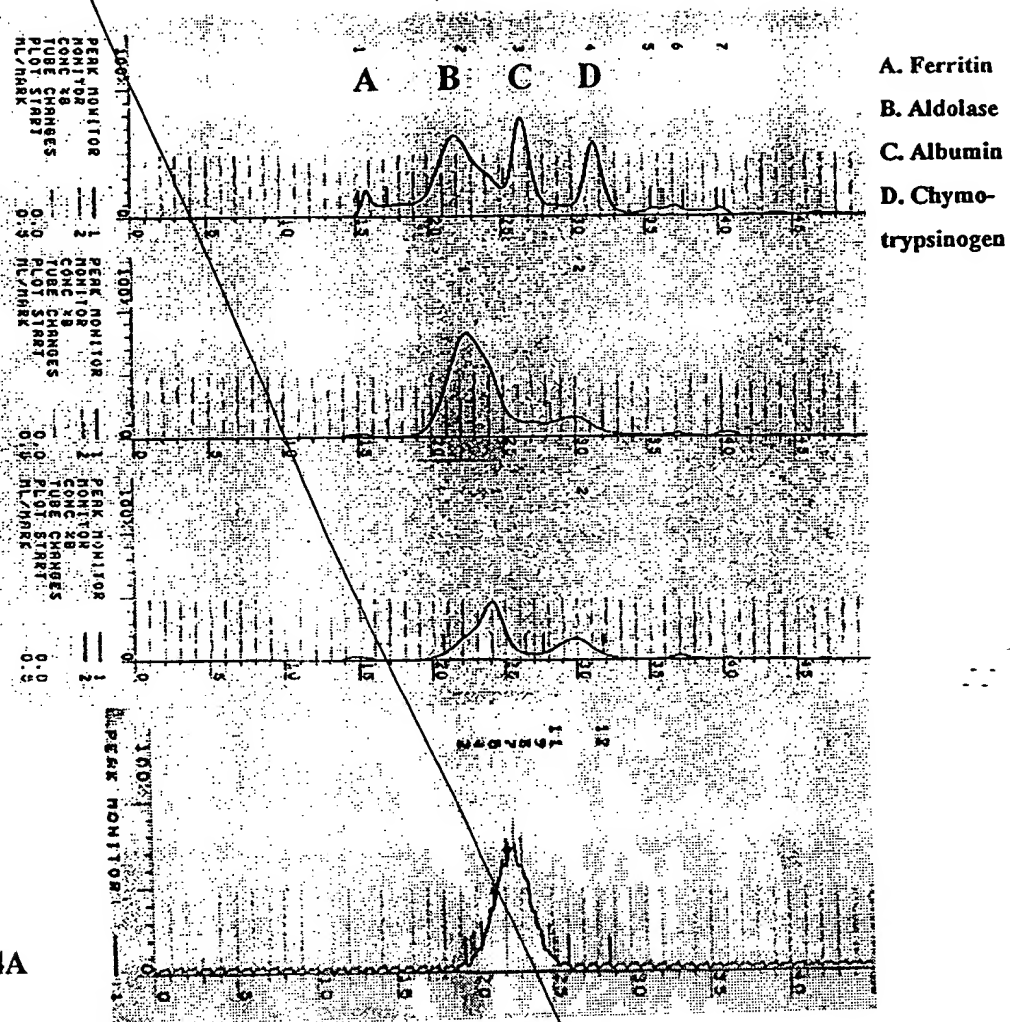


Corresponding U/ml/min by CBZ-Lys-S-Thiobenzyl Ester/DNTB

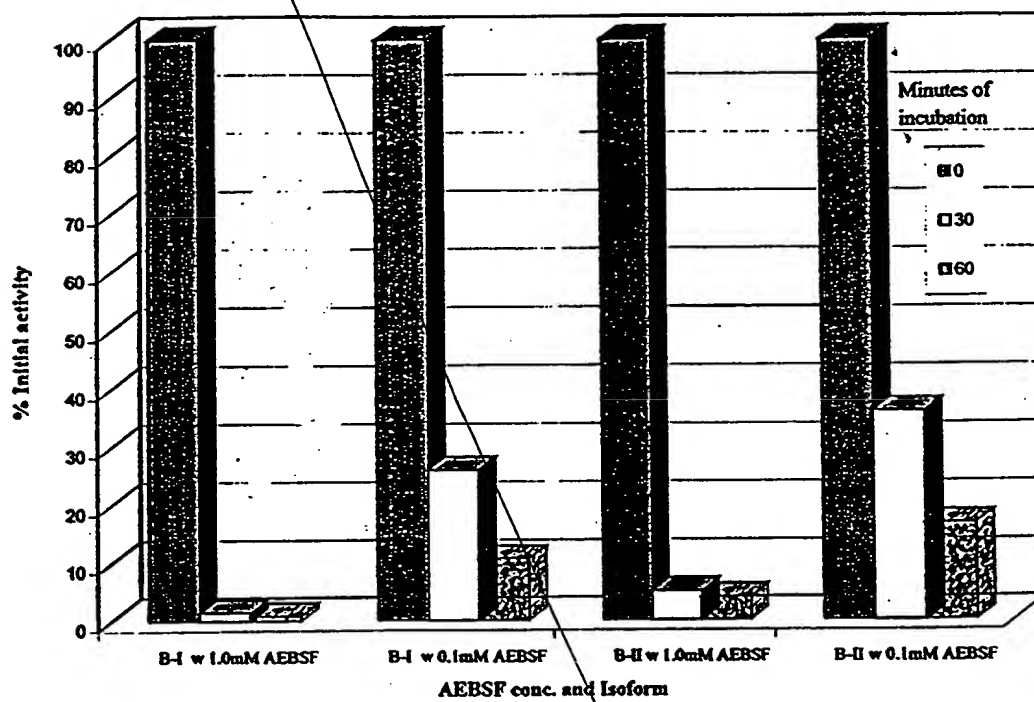
## Coupled cleavage Assay

**FIG. 6**

rh  $\beta$ -I  
tryptase  
Mutant S194A

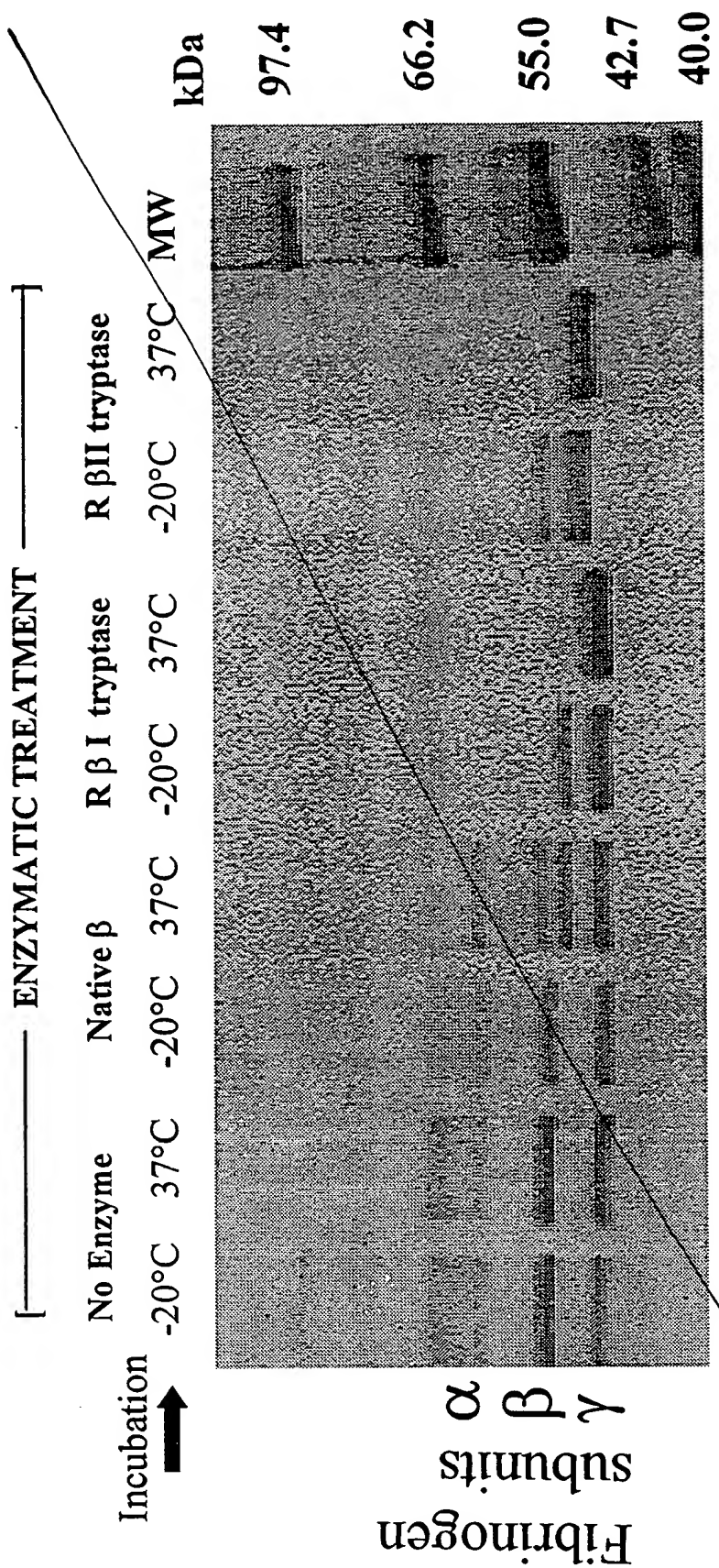


**FIG. 7**



**FIG. 8**





**FIG. 9**